Initial Approval: October 12, 2016

Revision Dates: January 11, 2017

## **CRITERIA FOR PRIOR AUTHORIZATION**

Tecentriq® (atezolizumab)

PROVIDER GROUP Professional

**MANUAL GUIDELINES** The following drug requires prior authorization:

Atezolizumab (Tecentriq®)

## **CRITERIA FOR APPROVAL FOR ATEZOLIZUMAB** (must meet all of the following):

- Patient must have one of the following:
  - Patient must have a diagnosis of locally advanced or metastatic urothelial carcinoma
    - Patient must have one of the following:
      - Disease progression during or following platinum-containing chemotherapy OR
      - Disease progression within 12 months of neoadjuvant or adjuvant treatment with platinum-containing chemotherapy
  - Patient must have a diagnosis of metastatic non-small cell lung cancer (NSCLC)
    - Patient must have disease progression during or following platinum-based chemotherapy
    - Patients with EGFR or ALK genomic tumor aberrations should have disease progression on FDA approved therapy for these aberrations prior to receiving Tecentriq.
- Must be prescribed by or in consultation with an oncologist or hematologist
- Patient must be 18 years of age or older
- Must be administered by a healthcare professional
- Patient must not be pregnant

**LENGTH OF APPROVAL:** 12 months

DATE

DRUG UTILIZATION REVIEW COMMITTEE CHAIR	PHARMACY PROGRAM MANAGER
	DIVISION OF HEALTH CARE FINANCE
	KANSAS DEPARTMENT OF HEALTH AND ENVIRONMENT

DATE